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CONCERNING A FILING UNDER 35 U.S.C. 371'INTERNATIONAL APPLICATION NO.
PCT/US99/05751INTERNATIONAL FILING DATE
3/16/1999PRIORITY DATE CLAIMED
3/16/1999

TITLE OF INVENTION RAPID DIAGNOSTIC METHOD FOR DISTINGUISHING ALLERGIES AND INFECTIONS

APPLICANT(S) FOR DO/EO/US UNIVERSITY OF FLORIDA, Parker Small, Shih-Wen Huang

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. ☒ This is a **FIRST** submission of items concerning a filing under 35 U.S.C. 371.
2. ☐ This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing under 35 U.S.C. 371.
3. ☐ This is an express request to begin national examination procedures (35 U.S.C. 371 (f)). The submission must include items (5), (6), (9) and (21) indicated below.
4. ☒ The US has been elected by the expiration of 19 months from the priority date (Article 31).
5. ☒ A copy of the International Application as filed (35 U.S.C. 371 (c)(2))
 - a. ☒ is attached hereto (required only if not communicated by the International Bureau).
 - b. ☒ has been communicated by the International Bureau.
 - c. ☒ is not required, as the application was filed in the United States Receiving Office (RO/US).
6. ☒ An English language translation of the International Application as filed (35 U.S.C. 371 (c)(2)).
 - a. ☒ is attached hereto.
 - b. ☒ has been previously submitted under 35 U.S.C. 154(d)(4).
- ☒ Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3))
 - a. ☐ are attached hereto (required only if not communicated by the International Bureau).
 - b. ☐ have been communicated by the International Bureau.
 - c. ☒ have not been made; however, the time limit for making such amendments has NOT expired.
 - d. ☐ have not been made and will not be made.
- ☐ An English language translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).
- ☐ An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)).
- ☐ An English language translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).

Items 11 to 20 below concern document(s) or information included:

11. ☐ An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
12. ☐ An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
13. ☐ A FIRST preliminary amendment.
14. ☐ A SECOND or SUBSEQUENT preliminary amendment.
15. ☐ A substitute specification.
16. ☐ A change of power of attorney and/or address letter.
17. ☐ A computer-readable form of the sequence listing in accordance with PCT Rule 13ter.2 and 35 U.S.C. 1.821 - 1.825.
18. ☐ A second copy of the published international application under 35 U.S.C. 154(d)(4).
19. ☐ A second copy of the English language translation of the international application under 35 U.S.C. 154(d)(4).
20. ☐ Other items or information:

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Timothy H. Van Dyke, Reg. No. 43,218

21. ☒ The following fees are submitted:**BASIC NATIONAL FEE (37 CFR 1.492 (a) (1) - (5)) :**Neither international preliminary examination fee (37 CFR 1.482)
nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO
and International Search Report not prepared by the EPO or JPO \$1000.00International preliminary examination fee (37 CFR 1.482) not paid to
USPTO but International Search Report prepared by the EPO or JPO \$860.00International preliminary examination fee (37 CFR 1.482) not paid to USPTO
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but all claims did not satisfy provisions of PCT Article 33(1)-(4) \$690.00International preliminary examination fee (37 CFR 1.482) paid to USPTO
and all claims satisfied provisions of PCT Article 33(1)-(4) \$100.00**ENTER APPROPRIATE BASIC FEE AMOUNT =**

\$ 100.00

Surcharge of \$130.00 for furnishing the oath or declaration later than ☐ 20 ☒ 30
months from the earliest claimed priority date (37 CFR 1.492(e)).

\$ 130.00

CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE	\$
Total claims	15	0	X \$18.00	\$ 0.00
Independent claims	3	0	X \$80.00	\$ 0.00
MULTIPLE DEPENDENT CLAIM(S) (if applicable)			+ \$270.00	\$ 0.00

TOTAL OF ABOVE CALCULATIONS =

\$ 230.00

☒ Applicant claims small entity status. See 37 CFR 1.27. The fees indicated above are
reduced by 1/2.

\$ 115.00

SUBTOTAL =

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NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR
1.137(a) or (b)) must be filed and granted to restore the application to pending status.

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RAPID DIAGNOSTIC METHOD FOR DISTINGUISHING ALLERGIES AND
INFECTIONS

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Background of the InventionI. Field of the Invention:

This invention is a rapid and simple method for the differential diagnosis of allergies,
10 sinusitis and upper respiratory tract infections. The method involves the use of either
commercially available or novel, specifically adapted, indicator or reagent test strips which
are contacted with nasal secretions. Based on the differential read-out from the indicator
strip, and a measure of eosinophil infiltration in the nasal secretion, a user of the strip is
able to determine, with the assistance of a scoring method disclosed herein, whether an
15 allergic condition or an infection is the cause of the respiratory discomfort.

II. Background:

It is common for patients afflicted with respiratory discomfort to seek the advice of a
20 clinician in an effort to minimize or overcome their discomfort. Such discomfort generally
is attributable to one of the following etiologies: allergic reactions, viral upper respiratory
tract infections (URIs), or bacterial infections which can produce sinusitis. However, the
clinician presented with such a patient typically has the daunting task of determining which
of these three principal etiologies is responsible for the discomfort experienced by a
25 particular patient. The danger
inherent in a mis-diagnosis can be quite severe. For example, should the clinician
incorrectly diagnose an allergy as sinusitis, a course of antibiotics would typically be
prescribed. Naturally, such treatment would do little to alleviate the allergic discomfort
being experienced by the patient while at the same time, the patient is exposed to an
30 antibiotic to which there is a possibility of raising a resistant bacterial infection. Should
this occur, a problem much more severe than the original allergic condition will have been
unwittingly engendered. The prevalence of antibiotic-resistant strains on a global scale
due to the over-prescription of antibiotics has become an increasingly recognized problem
(Service, R.F., 1995).

In the foregoing example, the availability of a rapid and simple differential diagnostic method would, instead of resulting in a compounded problem, result in the simple recommendation by the clinician that the patient adhere to a course of anti-histamine treatment, allergen avoidance, and/or a regimen of toleragenic desensitization. Unfortunately, however, to date, there is no such simple procedure which will provide the clinician with the necessary differential diagnosis. The accepted method of diagnosis for bacterial sinusitis is expensive radiologic imaging (typically X-ray or CT-scan) of the patients sinuses (see Katz et al., 1995).

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Many scientific articles have appeared addressing one or another of the various etiologies of respiratory discomfort. However, no rapid, inexpensive differential diagnostic method has been found. Thus, for example, Wang *et al.*, *Correlations between Complaints, Inflammatory Cells and Mediator Concentrations in Nasal Secretions after Nasal Allergen Challenge and during Natural Allergen Exposure*, Int. Arch. Allergy Immunology 1995; 106:278-285, disclosed a method of using a nasal microsuction technique. They showed that nasal allergen challenge (NAC) of asymptomatic (out of season) seasonal allergic rhinitis patients results in immediate (5 minutes) increases in histamine, leukotriene C4 (LTC4), and tryptase, with a more gradual (one hour post NAC) and prolonged increase in eosinophil and eosinophil cationic protein (ECP) concentration in nasal secretions. By contrast, in symptomatic (in season allergic rhinitis) patients, high concentrations of eosinophils, ECP, LTC4 and histamine, but not tryptase, were observed. It was concluded that allergic rhinitis is a chronic inflammation of the nasal mucosa, and that infiltration of eosinophils and release of late-phase inflammatory mediators are the predominant pathophysiologic markers. However, this publication neither teaches nor suggests that these observations can be applied to distinguish patients suffering from an allergic condition as opposed to an infection. In addition, the methods used by these authors are laborious and time-consuming and do not involve the use of reagent test strips. Sigurs *et al.*, *Eosinophil cationic protein in nasal secretion and in serum and myeloperoxidase in serum in respiratory syncytial virus bronchiolitis: relation to asthma and atopy*, Acta Paediatr 1994; 83:1151-5, concluded that it is not possible to predict, from eosinophil cationic protein/albumin ratios in nasal secretions or from ECP and myeloperoxidase concentrations in serum, whether children with respiratory syncytial virus (RSV) bronchiolitis would develop asthma. The publication provides no teaching or suggestion

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elected for publication

of a method which can easily distinguish patients suffering from an allergic condition as opposed to an infection. The methods used by these authors are laborious and time-consuming and do not involve the use of reagent test strips. Okuda *et al.*, *A Novel Method of Counting Eosinophils in Nasal Secretion of*

- 5 *Allergic Rhinitis by Hemocytometric Method*, Int. Arch. Allergy Immunol. 1994; 104 (suppl. 1):6, disclosed a rapid method for quantitating the number of eosinophils in nasal secretions as a method for diagnosis of allergic rhinitis. The method involves preparation of a solubilized sample of nasal secretion and counting of whole eosinophils. There is no mention of reagent test strips and there is no mention of a method for distinguishing
- 10 patients suffering from an allergic condition as opposed to an infection.

Kowalski *et al.*, *Neutrophil chemotactic activity (NCA) in nasal secretions from atopic and nonatopic subjects*, Allergy 1993; 48:409-414, reported that basal nasal secretions of both healthy persons and patients with chronic rhinitis contain significant chemotactic activity

15 to neutrophils. The study also reports that there is an increase in protein content of nasal secretions in patients with perennial allergic rhinitis (AR) following challenge with an antigen. There is no mention of reagent test strips and there is no mention of a method for distinguishing patients suffering from an allergic condition as opposed to an infection.

- 20 Igarashi *et al.*, *Analysis of nasal secretions during experimental rhinovirus upper respiratory infections*, J. Allergy Clin. Immunol. 1993; 92:722-731, studied patients with allergic rhinitis or control subjects inoculated with rhinovirus. Nasal lavage samples pre- and post-infection were analyzed for protein and mast cell mediators. It was found that total protein (including the plasma proteins albumin and IgG and the glandular proteins
- 25 lactoferrin, lysozyme and secretory IgA) increased post-infection, predominantly due to increased vascular permeability. It was also found that the allergic subjects had fewer symptoms, but greater vascular permeability and greater histamine secretion than control subjects post-rhinovirus infection. Protein was determined by the bicinchoninic acid protein assay (Pierce Chemical Co.) on aliquots of nasal lavage. It is noted at several
- 30 points in the publication that the symptoms of rhinovirus infected patients and patients with nasal allergic reactions are similar, thus teaching away from the possibility that a simple nasal secretion assay method could be used to distinguish these conditions. Inasmuch as this study is directed at determining the differences in nasal secretions between rhinovirus infected normal or allergic individuals, the study addresses a different

problem than that addressed by the instant invention, which is a method for measuring the differences in nasal secretions of patients infected with a rhinovirus, for example, and a patient not infected with a rhinovirus but suffering from allergic rhinitis. The methods used by these authors are laborious and time-consuming and do not involve the use of reagent test strips.

Sperber *et al.*, *In vivo detection of a novel macrophage-derived protein involved in the regulation of nasal mucus-like glycoconjugate secretion*, J. Allergy Clin. Immunol. 1993; 92:581-588, disclosed a study directed at characterization of a novel 68 Kd nasal mucus secretagogue (NMS-68) released by monocytes. Inasmuch as this protein was found to be present in nasal tissue of patients with allergic and non-allergic rhinitis, it does not appear to provide a method for distinguishing between these conditions (although the baseline level of this protein is more elevated in the allergic patients). Reagent test strips were not used in this study.

Knani *et al.*, *Indirect evidence of nasal inflammation assessed by titration of inflammatory mediators and enumeration of cells in nasal secretions of patients with chronic rhinitis*, J. Allergy Clin. Immunol. 1992; 90:880-889, examined the nasal lavage cells and six inflammatory mediators released in nasal secretions of four groups of patients with perennial rhinitis and a control group. It was found that patients with symptomatic allergic rhinitis had increased levels of eosinophils, as well as of eosinophil protein X (EPX), LTC₄/D₄, tryptase, MPO and PGD₂. Patients with non-allergic rhinitis were found to have increased neutrophil, tryptase, MPO and EPX concentrations. These measurements provided indirect evidence of nasal inflammation. However, there was no analysis of differences between the nasal secretions of rhinitis patients and patients suffering, for example, from a rhinoviral infection or a bacterial sinus infection.

Klementsson *et al.*, *Eosinophils, secretory responsiveness and glucocorticoid-induced effects on the nasal mucosa during a weak pollen season*, Clinical and Experimental Allergy 1991; 21:705-710, analyzed the eosinophil influx, the concentration of ECP and secretory responsiveness following methacholine challenge in nasal lavage samples of patients with allergic rhinitis. There was no concurrent analysis of nasal secretions from patients with bacterial sinus or viral infections, and reagent test strips were not employed.

Gordon *et al.*, *The pathophysiology of rhinitis*, J. Allergy Clin. Immunol. 1991; 88:33-42. challenged patients with seasonal rhinitis on one side of the nose with an allergen and nasal secretions from both sides of the nose were analyzed for protein and mediators. There was no concurrent analysis of nasal secretions from patients with bacterial or viral infections. and reagent test strips were not employed.

Cohen, R. A. and Brestel, E. P., *Nasal secretory response to allergen provocation*, 1988; 18:435-443, analyzed nasal lavage samples from ragweed-sensitive and control subjects following ragweed pollen challenge. The study revealed no increase in total protein, albumin, potassium, lysozyme activity or peroxidase activity in the control subjects. There were increases in all of these constituents in the ragweed-sensitive subjects. These constituents were assayed by dye binding (Bradford), rocket immunoelectrophoresis, absorption spectroscopy, ABTS (Sigma Chemical Co.) oxidation, and radial diffusion, respectively. Use of reagent test strips is neither taught nor suggested, nor is there a concomitant analysis of these constituents in bacterially or viral infected, no-allergic subjects.

Liu *et al.*, *Injurious effect of eosinophil extract on the human nasal mucosa*, Rhinology 1988; 26:121-132, attempted to elucidate the role of eosinophils in nasal secretions of allergic subjects. These authors conclude that the eosinophil extracts tested may actually be harmful to the function of human nasal mucosa. Use of reagent test strips is neither taught nor suggested, nor is there a concomitant analysis of these constituents in infected subjects.

Anderson *et al.*, *Allergen-induced nasal hyperactivity appears unrelated to the size of the nasal and dermal immediate allergic reaction*, Allergy 1987, 42:631-637, analyzed "priming", in which nasal lavage samples from hay fever patients were tested following an initial and a re-challenge with allergen. The biochemical parameter used as the measure of the allergic reaction was TAME-esterase via a radiochemical method (release of tritium labeled methanol from the synthetic substrate H3-TAME). Use of reagent test strips is neither taught nor suggested, nor is there a concomitant analysis of these constituents in infected subjects.

Settipane, G. A. and Klein, D. E., *Non Allergic Rhinitis: Demography of Eosinophils in Nasal Smear, Blood Total Eosinophil Counts and IgE Levels*. NER Allergy Proc.1985; 6:363-366. in an attempt to develop a methodology for differential diagnosis of patients with non-allergic rhinitis, evaluated patients with rhinitis and negative skin tests, (taken to mean that their rhinitis had a non-allergic etiology), for cause of the rhinitis. Nasal smears from these patients were obtained by rolling a swab with nasal secretions on a glass slide, fixing with methanol, staining with Camaco stain (Wright-Giemsa stain), and counting the number of eosinophils per 100 cells. Sinus X-rays were conducted to detect sinusitis. NARES, non-allergic rhinitis with eosinophilia syndrome, is clinically defined in the paper as "nasal congestion/rhinorrhea with negative allergy skin tests, normal serum IgE, and $\geq 5\%$ eosinophils in the nasal smear." The purpose of the study was "to attempt to corroborate NARES as a new syndrome and to attempt to further classify and clarify non allergic rhinitis." There was no discussion in this paper regarding the problem of misidentification of this clinical condition with upper respiratory tract infections, and use of reagent test strips as part of the differential diagnosis is neither disclosed nor suggested.

Brofeldt, *et al.*, *Biochemical Analysis of Nasal Secretions induced by Methacholine, Histamine, and Allergen Provocations*, Am. Rev. Respir. Dis. 1986; 133:1138-1142. obtained methacholine, histamine and allergen induced nasal secretions from subjects over a fifteen minute post-induction period. The nasal secretions were weighed and tested for hexose content (orcinol method), protein content (Lowry method), carbohydrate (gas liquid chromatography), sialic acid (colorimetric thiobarbituric acid assay), inorganic sulphate (radioactive BaCl₂), DNA (diphenylamine), albumin and immunoglobulins (rocket immunoelectrophoresis or ELISA). The different inducers were found to have differential effects on the various elements tested. However, there was no concomitant study of nasal secretions from patients suffering from an infection nor was the use of reagent test strips taught or suggested.

Eggelston, *et al.*, *Mediators of Immediate Hypersensitivity in Nasal Secretions during Natural Colds and Rhinovirus Infection* Acta Otolaryngol. 1984; suppl. 413:25- 35, note that "[v]iral respiratory infections and allergic rhinitis have many similarities. Not only are symptoms similar in the two conditions, but the pathologic anatomy of both is dominated by vascular dilatation and edema with minimal cellular infiltrate in acute phases ..." The

authors postulated that these similarities are due to mast cell activation during infection. resulting in release of histamine. However, they report that spectrofluorometric analysis of histamine in nasal secretions of control subjects or patients with a natural cold or with a rhinovirus infection does not support this hypothesis. Histamine concentrations were
5 found to be generally lower in individuals infected with influenza A or rhinovirus. TAME-esterase was also found not to be elevated during these infections. This article reports that during viral respiratory tract infection there is little or no esterase elevation, while in allergic rhinitis there is esterase elevation (see discussion at page 34 of the reference). However, the use of reagent test strips for this purpose is neither taught nor suggested. nor
10 is there a discussion of how these results could be used in a differential diagnostic method also aimed at distinguishing sinusitis.

Baumgarten. *et al*, *Plasma Kallikrein During Experimentally Induced Allergic Rhinitis: Role in Kinin Formation and Contribution to TAME-Esterase Activity in Nasal Secretions*.
15 *J. Immunol.* 1986; 137:977-982, report the observation that when allergic and non-allergic individuals are challenged intranasally with an allergen, post-challenge nasal lavages of only the allergic individuals contained elevated levels of immunoreactive human plasma kallikrein/prekallikrein (iHPK). This increase in iHPK correlated with increases in kinins, histamine, TAME-esterase, and clinical symptoms. In fact, these researchers argue that the
20 TAME-esterase activity is produced by plasma kallikrein and mast cell tryptase. Thus, iHPK may be an additional marker that could be used in a reagent test strip for allergic rhinitis diagnosis. However, in this study, esterase was assayed by a radiochemical method and kinins were assayed using a radioimmunoassay, rather than by any type of reagent test strip.

25 Anderson *et al*, *Mechanisms of nasal hyper-reactivity*. *Eur. Arch. Otorhinolaryngol.* 1995; 252 (suppl. 1):S22-S26, review the factors known to be involved in allergen-induced nasal hyper-reactivity. However, there is no teaching or suggestion of a method for distinguishing allergic from infection related rhinitis. Reagent test strips for this purpose
30 are neither proposed nor suggested.

Florman, *et al.*, *Rapid Non-invasive Techniques for Determining Etiology of Bronchitis and Pneumonia in Infants and Children*. *Clin. Chest Med.* 1987; 8:669-679, provide a review of techniques for differential diagnosis of the causative agent in lower respiratory

tract infections. A number of rapid, non-specific and specific tests are mentioned. However, there is no mention of differential diagnosis of allergic from infective conditions and reagent test strips for this purpose are neither disclosed nor suggested.

- 5 Katz *et al.*, *A comparison of Imaging Techniques in Patients with Chronic Sinusitis* (X-Ray, MRI, A-Mode Ultrasound) *Allergy Proc.* 1995. 16:123-127, demonstrate the long-felt need for a rapid, inexpensive way to diagnose sinusitis and distinguish this condition from allergic rhinitis. The method of the instant invention has the potential for supplanting the much more expensive diagnostic techniques reported in this paper to be most reliable in
10 diagnosis of sinusitis (CAT scans and MRI).

Demoly *et al.*, *Assessment of Inflammation in noninfectious chronic maxillary sinusitis*, *J. Allergy Clin. Immunol.* 1994; 94:95-108, suggest that it might be possible to distinguish sinusitis (infection) from allergic rhinitis based on the contents of nasal mucosa. However,
15 the techniques used in attempting to distinguish these conditions depended on the use of immunohistochemistry of surgical specimens, immunocytochemistry of lavage fluids, and measurement of specific inflammatory mediators in sinus lavage fluids (ELISA, RIA). There is no teaching or suggestion that reagent dip-sticks could be used for this purpose.

- 20 Accordingly, there has been a long-felt need in the art for a rapid, inexpensive, non-invasive technique for a method capable of distinguishing between allergies and infections. The method of the instant invention involves testing nasal secretions with commercially available (Ames Division, Miles Laboratories, Inc., Elkhart, IN 46515) or novel or modified reagent test strips. The commercially available strips, also referred to as
25 dipsticks, test for pH, protein, glucose, ketone, white blood cell esterase, bilirubin and blood.

The following U.S. Patents which may be relevant to the instant invention are listed on the product insert of the Miles Laboratories Inc., reagent test strips, and are herein
30 incorporated by reference:

- 1) 3,438,737- Protein test composition and method of detecting proteins in fluids using the test composition.
- 2) 4,301,115- Test device having resistance to cross contamination between reactant areas and process for making it.

- 3) 4,637,979- Composition and test device for determining the presence of leukocytes containing a zwitterion coupling agent for determining the presence of leukocytes, esterase or protease in a sample.
- 4) 4,645,842- Pyrrole composition for detecting presence of hydrolytic analytes, useful in the detection of leukocytes, esterase and protease in a test sample.
- 5) 4,657,855- Composition and test device for determining the presence of leukocytes, esterase and protease in a test sample.
- 6) 4,704,460- Novel compounds for detecting the presence of hydrolytic analytes in a test sample which indicates the presence of leukocytes, esterase and protease in a test sample.
- 7) 4,758,508- Analytical process and agents for the detection of esterolytic and/or proteolytic enzymes in a liquid sample.

The following U.S. Patents which may be relevant to the instant invention are listed on the product insert or packaging of Boehringer Mannheim Corporation's reagent test strips, and are herein incorporated by reference:

- 1) 3,359,072- A protein determination method.
- 2) 3,418,079- A protein determination device and method.
- 3) 3,712,853- A nitrite detection reagent and method.
- 4) 3,897,214- A diagnostic device.
- 5) 3,802,842- A reagent test strip.
- 6) 4,013,416- A protein detection method.
- 7) 4,385,114- An oxidation indicator system.

However, none of these patents disclose or suggest a method for testing nasal mucous secretions to distinguish allergic from infectious conditions.

In the method of this invention, a sample of a patient's nasal secretions is tested and, based on the pH, amount of protein, nitrite and leukocyte esterase and eosinophils, it can quickly be determined if the patient is suffering from an allergy, a simple viral infection, or bacterial sinusitis. The method has the potential to supplant much more expensive and invasive clinical procedures.

Brief Summary of the Invention

This invention provides a method for rapidly, non-invasively and inexpensively differentiating between allergic rhinitis and respiratory viral or bacterial infections. The method involves measuring, for example by contacting a reagent test strip with a sample of nasal secretion, a series of agents in nasal secretions. The reagent test strip provides information on the pH, protein content, nitrite content, esterase activity and preferably also provides information on the level of eosinophil infiltration in the sample contacted, such that a combination of a pH between about 7.5 and 9, a moderately strong presence of protein, at least a trace of esterase and nitrite and little or no eosinophil infiltration indicates the presence of a bacterial infection (sinusitis). The combination of a pH between about 5.0 and 7.5, little or no protein, little or no esterase activity and little or no eosinophil infiltration, is an indication of viral infection (URI). However, the same profile as in URI viral infection but with clear indication of eosinophil infiltration is an indication of allergic rhinitis.

Also provided is a reagent test strip specifically adapted for rapidly, non-invasively and inexpensively differentiating between allergy and respiratory infection. The reagent test strip is adapted to provide information on the pH, protein content, nitrite content, esterase activity and eosinophil infiltration of the sample contacted, such that the method of this invention can quickly and easily be practiced.

Brief Summary of the Figures

Figure 1 shows the different chemical read-outs obtained by contacting reagent test strips with the nasal secretions of multiple patients presenting with respiratory discomfort. Closed circles (●) represent patients with radiologically proven sinusitis; open circles (○) represent patients having no sinusitis based on radiological investigation; open squares (□) represent patients with radiologically proven sinusitis that were at least partially treated with antibiotics prior to or during the evaluation period.

Figure 2 is a histogram based on the data shown in Figure 1 and a measure of eosinophil infiltration in the nasal exudates of the Figure 1 patients (not shown in Figure 1). From the raw data of Figure 1, a point system was established such that strong evidence of

eosinophil infiltration (+ + + eosinophil) received negative score of -6, moderate infiltration (+ + eosinophils) received a score of -4, and trace infiltration (+ eosinophils) received a score of -2; pHs greater than or equal to 8.5 received a score of +3, a pH between 8.0 and 8.4 received a score of +2, a pH between 7.25 and 7.9 received a score of +1 and a pH between 5.0 and 7.25 received a score of zero; protein content between about 30-100 mg/dl was given a score of +1; between about 100 mg/dl and 300 mg/dl was given a score of +2; protein between about 300 mg/dl and 2000 mg/dl was given a score of +3; and protein greater than 2000 mg/dl received a score of +4; low nitrite concentrations received a score of +1; moderate amounts of nitrite received a score of +2 and high amounts of nitrite received a score of +3; esterase activity in small amounts received a score of about +1, a score of +2 was assigned moderate amounts and a score of +3 was assigned to large amounts of esterase. By summing the scores assigned to eosinophils, pH, nitrite, protein, and esterase, the histogram of Figure 2 was produced. The level of eosinophil infiltration is shown in the figure by + , + +, and + + + for trace, moderate and strong eosinophil infiltration, respectively. Patients radiologically confirmed to have sinusitis are shown with stippled symbols; patients radiologically confirmed to have sinusitis but who had been at least partially treated with antibiotics are shown with a half filled half stippled symbol; patients radiologically clear of sinusitis but which exhibited eosinophilia are shown by open symbols; patients radiologically clear of sinusitis without concomitant eosinophilia are shown with diagonally filled/empty symbols.

Detailed Description of the Invention

This invention is a method for non-invasively, rapidly and simply distinguishing between allergies and infections which involves testing nasal secretions for the levels of a number of agents. Preferably the method comprises contacting nasal secretions with commercially available (for example, from Ames Division, Miles Laboratories, Inc., Elkhart, IN 46515, or from Boehringer Mannheim Corporation, Advanced Diagnostics, 9115 Hague Road, P.O. Box 50457, Indianapolis, IN 46250-0457) or novel specifically adapted reagent test strips. The commercially available strips, also referred to as dipsticks, typically test for pH, protein, nitrite glucose, ketone, white blood cell esterase, bilirubin and blood. In the method of this invention, a sample of a patient's nasal secretions is contacted with a reagent test-strip and, based on the pH, presence or absence of protein, levels of nitrite, esterase and a measure of eosinophil infiltration in the nasal secretion, it can quickly be determined

if the patient is suffering from an allergy or an infection. The method has the potential to supplant much more expensive and invasive clinical procedures. In the method of this invention, a commercially available reagent test-strip, or a novel, specifically adapted reagent test-strip, is contacted with a sample of a patient's nasal secretions. The patient
5 may be any mammal, including an animal or a human. Based on the differential read-out from the reagent test-strip, which preferably includes a measure of eosinophil infiltration, a determination can be made as to whether an allergy or an infection is responsible for upper-respiratory tract discomfort. With respect to commercially available reagent test-strips, those sold by the Ames Division of Miles Laboratories or the Boehringer Mannheim
10 Corporation are generally acceptable. These test-strips may be made as disclosed herein or as disclosed in any of U.S. Patent Nos. 3,438, 737, 4,301,115; 4,637,979, 4,645,842; 4,657,855; 4, 704,460; 4, 758,508; 3,359,072; 3,418,079, 3, 712,853; 3,897,214; 3,802,842; 4,013,416; and 4,385,114; all of which are incorporated by reference herein for this purpose. These reagent test-strips may be employed in the novel method of this
15 invention as further described hereinbelow with the additional requirement being that a measure of eosinophils must be made.

Accordingly, one embodiment of this invention is a simple method for distinguishing patients with allergies and patients with upper respiratory infections involves testing the
20 nasal secretions of these patients with reagent strips disclosed in the aforementioned patents. These dipsticks are inexpensive, and provide an indication of a contacted fluid's pH, protein, glucose, ketone, nitrite, esterase, bilirubin and blood content. The patient blows his/her nose on a receptacle (for
example wax paper or plastic film such as Saran Wrap or the like), or a swab of nasal
25 secretion is taken, and contacted with the test strip. The pH, protein, nitrite, and esterase contents are then evaluated based on the directions of the manufacturer found on the outside of the box in which the commercially available reagent test strips are sold. The procedure generally takes less than about sixty seconds. The following table provides results and the meaning thereof:

	Allergy	Viral Infection	Bacterial Infection (Sinusitis)
pH	5.0-7.5	7.5-9	7.5-9
Protein	Trace to +	Trace to +	++ or +++
Nitrite	Negative or trace	Negative or trace	++ or +++
Esterase	Negative or trace	Trace to +	++ or +++
Eosinophils	++ or +++	- or trace	- or trace

This data can be visualized by referring to Figure 1 which provides a graphic representation of the different chemical read-outs obtained by contacting reagent test strips with the nasal secretions of multiple patients presenting with respiratory discomfort.

- 5 Closed circles (●) represent patients with radiologically proven sinusitis; open circles (○) represent patients having no sinusitis based on radiological investigation; open squares (□) represent patients with radiologically proven sinusitis that were at least partially treated with antibiotics prior to or during the evaluation period. As can be seen from Figure 1, there is a clustering in the data points for the pH, protein, nitrite and esterase contents of
- 10 patients with or without sinusitis.

To aid in the differential diagnosis of allergic rhinitis, viral URI and sinusitis, we have discovered that by assigning a point system to several parameters measured from the mucus secretions of the patients, delimited differential scattering of the patient populations

15 can be achieved with only a minor amount of overlap between patients having these various clinical conditions. Figure 2 is a histogram based on the data shown in Figure 1 and a measure of eosinophil infiltration in the nasal exudates of the Figure 1 patients (not shown in Figure 1). From the raw data of Figure 1, a point system was established such that strong evidence of eosinophil infiltration (+++ eosinophil) received negative score of

20 -6, moderate infiltration (++ eosinophils) received a score of -4, and trace infiltration (+ eosinophils) received a score of -2; pHs greater than or equal to 8.5 received a score of +3, a pH between 8.0 and 8.4 received a score of +2, a pH between 7.25 and 7.90 received a score of +1 and a pH between 5.0 and 7.25 received a score of zero; protein content between about 30-100 mg/dl was given a score of +1; between about 100 mg/dl and 300

25 mg/dl was given a score of +2; protein between about 300 mg/dl and 2000 mg/dl was given a score of +3 and protein greater than 2000 mg/dl received a score of +4; low nitrite concentrations received a score of +1; moderate nitrite concentrations received a score of

+2; high concentrations of nitrite received a score of +3; esterase activity in small amounts received a score of about +1, a score of +2 was assigned to moderate amounts and a score of +3 was assigned to large amounts of esterase. By summing the scores of the eosinophils, pH, nitrite, protein, and esterase, the histogram of Figure 2 was produced. The level of eosinophil infiltration is shown in the figure by +, ++, and +++ for trace, moderate and strong eosinophil infiltration, respectively. Patients radiologically confirmed to have sinusitis are shown with stippled symbols; patients radiologically confirmed to have sinusitis but who had been at least partially treated with antibiotics are shown with a half filled half stippled symbol; patients radiologically clear of sinusitis but which exhibited eosinophilia are shown by open symbols; patients radiologically clear of sinusitis without concomitant eosinophilia are shown with diagonally filled/empty symbols. This method of differential diagnosis can be used with any method of collecting the pH, protein, esterase, nitrite and eosinophil content of nasal secretions, and is not restricted to the use of reagent indicator strips. The use of reagent indicator strips is, however, one of the most easily conducted, inexpensive and rapid methods for achieving this analysis.

The level of eosinophil infiltration is easily estimated in one embodiment of the invention using high power field microscopy (about 400 x magnification) by determining the percentage of total cells that are eosinophils in a Hansel's stained smear of nasal secretion. Accordingly, a 0-10% eosinophil profile received no negative score: a 10-25% eosinophil profile received a "+" (i.e. -2 assigned score); a 26-50% eosinophil profile received a "++" (i.e. -4 assigned score); and a 51-75% eosinophil profile received a "+++" (i.e. an assigned score of -6).

According to the package insert of the AMES 9SG Multistix® product, the following information, supplied for urinalysis by that manufacturer, may be directly applied to the novel utility disclosed and claimed herein, and is hereby incorporated as follows:

REAGENT STRIPS for mucus analysis are firm plastic strips to which are affixed several separate reagent areas. Depending on the product being used, AMES REAGENT STRIPS provide tests for glucose, bilirubin, ketone (acetoacetic acid), specific gravity, blood, pH, protein, urobilinogen, nitrite, and leukocytes in urine. A user of these strips refers to the carton and bottle label for specific reagent areas on the product being used.

The reagent test areas on AMES REAGENT STRIPS are ready to use upon removal from the bottle and the entire reagent strip is disposable. The strips may be read visually, requiring no additional laboratory equipment for testing. Certain configurations of strips may also be read instrumentally, using the CLINITEK® family of Urine Chemistry

5 Analyzers and the appropriate Program Module or Program Card from AMES.

The directions must be followed exactly. Accurate timing is essential to provide optimal results. The reagent strips must be kept in the bottle with the cap tightly closed to maintain reagent reactivity. To obtain optimal results, it is necessary to use fresh mucus.

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Chemical Principles of the Procedure:

pH: This test is based on a double indicator principle that gives a broad range of colors covering the entire urinary pH range. Colors range from orange through yellow and green
15 to blue.

Protein: This test is based on the protein-error-of-indicators principle. At a constant pH, the development of any green color is due to the presence of protein. Colors range from yellow for "Negative" through yellow-green and green to green-blue for "Positive"
20 reactions.

Nitrite: At the acid pH of the reagent area, nitrite in the nasal secretion reacts with p-arsanilic acid to form a diazonium compound. This diazonium compound in turn couples with 1,2,3,4-tetrahydrobenzo(h)quinolin-3-ol to produce a pink color. The intensity of the
25 pink color developed is used as the basis for assigning the nitrite concentration score as described above.

Leukocytes: Granulocytic leukocytes contain esterases that catalyze the hydrolysis of the derivatized pyrrole amino acid ester to liberate 3-hydroxy-5-phenyl pyrrole. This pyrrole
30 then reacts with a diazonium salt to produce a purple product. The intensity of the purple color developed is used to assign a value to esterase activity as described above.

REAGENTS (Based on dry weight at time of impregnation):

pH: 0.2% w/w methyl red; 2.8% w/w methyl red; 2.8% w/w bromthymol blue; 97.0% w/w non-reactive ingredients.

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Protein: 0.3% w/w tetrabromphenol blue; 97.3% w/w buffer; 2.4% w/w nonreactive ingredients.

Nitrite: 1.4% w/w p-arsanilic acid; 1.3% w/w 1,2,3,4-tetrahydrobenzo-(h)-quinolin-3-ol;
10 10.8% w/w buffer; 86.5% w/w nonreactive ingredients.

Leukocytes: 0.4% w/w derivatized pyrrole amino acid ester; 0.2% w/w diazonium salt;
40.9% w/w buffer; 58.5% w/w nonreactive ingredients.

15 Those skilled in the art will recognize that other chemical components may be used for carrying out the method disclosed herein.

Recommended Procedures for Handling Reagent Strips: All unused strips must remain in the original bottle. Transfer to any other container may cause reagent strips to deteriorate
20 and become unreactive. Do not remove desiccant(s) from bottle. Do not remove strip from the bottle until immediately before it is to be used for testing. Replace cap immediately and tightly after removing reagent strip. Do not touch test areas of the reagent strip. Work areas and specimen containers should be free of detergents and other contaminating substances.

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Dip test areas in mucus completely, but briefly, to avoid dissolving out the reagents. If using strips visually, read test results carefully at the times specified, in a good light (such as fluorescent) and with the test area held near the appropriate Color Chart on the bottle label. Do not read the strips in direct sunlight. If the strips are used instrumentally,
30 carefully follow the directions given in the appropriate instrument operating manual. Protection against ambient moisture, light and heat is essential to guard against altered reagent.

- Discoloration or darkening of reagent areas may indicate deterioration. If this is evident, or if test results are questionable or inconsistent with expected finding, the following steps are recommended: (1) confirm that the product is within the expiration date shown on the label; (2) check performance against known positive control materials (e.g., CHEK-
5 STIX® Control Strips); (3) retest with fresh product.

The following procedure should be followed exactly to achieve reliable test results:

1. Collect fresh mucus specimen on a non-absorbent surface.
2. Remove one strip from the bottle and replace cap. Completely immerse reagent
10 areas of the strip in fresh mucus and remove immediately to avoid dissolving out reagents.
3. If reading visually, compare reagent areas to corresponding Color Chart on the bottle label at the time specified. Hold strip close to color blocks and match carefully. Avoid laying the strip directly on the Color Chart, as this will result in
15 soiling the chart.
4. If reading instrumentally, carefully follow the directions given in the appropriate instrument operating manual.

Proper read time is critical for optimal results. If using strips visually, read the pH, protein, and nitrite at 60 seconds; and leukocytes at 2 minutes. The pH and protein areas may also
20 be read immediately or at any time up to 2 minutes after dipping.

After dipping the strip, check the pH area. If the color on the pad is not uniform, read the reagent area immediately, comparing the darkest color to the appropriate Color Chart. All reagent areas except leukocytes may be read at about 1 minute for identifying negative
25 specimens and for determination of the pH. A positive reaction (small or greater) at less than 2 minutes on the leukocyte test may be regarded a positive indication of leukocytes. Color changes that occur after 2 minutes are of no diagnostic value. If using strips instrumentally, the instrument will automatically read each reagent area at a specified time.

- 30 For best results, performance of reagent strips should be confirmed by testing known negative and positive specimens or controls whenever a new bottle is first opened. Negative and positive specimens or controls may also be randomly hidden in each batch of specimens tested. Each laboratory should establish its own goals for adequate standards of performance, and should question handling and testing procedures if these standards are

not met. CHEK-STIX® Urinalysis Control Strips (#1360) from AMES®, with positive or defined results, provide a convenient basis for quality control program.

Results with AMES REAGENT STRIPS are obtained in clinically meaningful units directly from the Color Chart comparison when using strips visually. With instrumental use, the reagent pads are "read" by the instrument and the results are displayed or printed. The color blocks and instrumental display values represent nominal values; actual values will vary around the nominal values.

10 pH: If proper procedure is not followed and excess mucus remains on the strip, a phenomenon known as "runover" may occur, in which the acid buffer from the protein reagent will run onto the pH area, causing a false lowering of the pH result.

Protein: Contamination of the nasal secretion specimen with quaternary ammonium compounds (e.g., from some antiseptics and detergents) or with skin cleansers containing chlorhexidine may also produce false positive results.

Nitrite: Pink spots or pink edges should not be interpreted as a positive result. Any degree of uniform pink color development should be interpreted as a positive nitrite test.

20 Leukocytes: Elevated glucose concentrations (≥ 3 g/dl) or high specific gravity may cause decreased test results. The presence of cephalexin (Keflex®), cephalothin (Keflin®), or high concentrations of oxalic acid may also cause decreased test results. Tetracycline may cause decreased reactivity, and high levels of the drug may cause a false negative reaction.

25 Leukocytes: Normal nasal secretion will generally yield negative results; positive results (small or greater) are clinically significant. Individually observed trace results may be of questionable clinical significance; however, trace results observed repeatedly may be clinically significant. Positive and repeated trace results indicate the need for further testing of the patient and/or nasal specimen, according to medically accepted procedures.

Specific Performance Characteristics: Specific performance characteristics are based on clinical and analytical studies. In clinical specimens, the sensitivity depends upon several factors: the variability of color perception; the presence or absence of inhibitory factors,

the specific gravity, and the pH; and the lighting conditions when the product is read visually. Because the color of each reagent area changes as the analyte concentration increases, the percentage of specimens detected as positive will increase with the analyte concentration.

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Each color block or instrumental display value represents a range of values. Because of specimen and reading variability, specimens with analyte concentrations that fall between nominal levels may give results at either level. Exact agreement between visual results and instrumental results might not be found because of the inherent differences between the

10 perception of the human eye and the optical system of the instruments.

pH: The pH test area measure pH values generally to within 1 unit in the range of 5-8.5 visually and 5-9 instrumentally.

15 Protein: The reagent area is more sensitive to albumin than to globulins, hemoglobin, Bence-Jones Protein and mucoprotein; a negative result does not rule out the presence of these other proteins.

Nitrite: Comparison of the reacted reagent area against a white background may aid in the
20 detection of low levels of nitrite ion, which may otherwise be missed.

Availability: AMES REAGENT STRIPS for Urinalysis are available in bottles of 100 strips: MULTISTIX® 10 SG (#2300A); MULTISTIX® 9 (#2301A); MULTISTIX® 9 SG (#2303A); MULTISTIX® 8 SG (#2304A); MULTISTIX® 7 (#2305A); N-MULTISTIX®
25 SG (#2740A); MULTISTIX® SG (#2741A); N-MULTISTIX® (#2829A); MULTISTIX® (#2820A); and BILI-LABSTIX® (#2814A).

Any of these or other commercially available reagent test strips which provide pH, protein, nitrite, esterase and preferably also eosinophil data can be used according to this disclosure
30 to differentiate between bacterial infections, viral infections and allergic conditions. Thus, in a fashion completely analogous to that described above for the Ames REAGENT STRIPS, commercially available reagent test strips produced by Boehringer Mannheim Corporation may be used or adapted for this purpose. For example CHEMSTRIP 9, Catalog No. 417109, provides a readout for leukocytes, nitrite, pH, protein and several

other analytes. The information provided in the package insert for the CHEMSTRIP 6, 7, 8, 9, 10 (which also provides a readout for specific gravity), is largely analogous to the information provided hereinabove from the Multistix® product. In our hands, testing of nasal secretions using the Boehringer product yielded results which, according to this invention, are similar to those obtained using the Multistix® product. Slight adjustments in the color readouts and values thereof may be needed due to the differences between the color charts used by the two manufacturers, but, based on the instant disclosure, those skilled in the art are able to make any necessary adjustments.

10 In one aspect of the invention, there is provided a reagent test strip specifically adapted for rapidly, non-invasively and inexpensively differentiating between allergic conditions and respiratory, viral and bacterial infections. In use, the test strip is optimized to provide information about nasal mucus secretions including, but not limited to, the pH, protein content, nitrite, esterase activity and level of eosinophil infiltration, such that all of the information presented in Figure 2 can be obtained from a single indicator strip. This is achieved, for example, by preparing a reagent test strip according to commercially available strips, but in addition, providing a means for measuring the amount of eosinophil cationic protein (ECP) or another eosinophil specific protein or enzyme present in the nasal secretion, the presence of which is proportional to the amount of eosinophil infiltration.

20 Alternatively, the test strip could be made so as to selectively trap eosinophils, and the assay could then be for any substance found in eosinophils (such as an enzyme or any other detectable substance), without it being necessary for the substance to be specific for eosinophils. The novel reagent test strip of this invention, therefore, can include an indicator location on the strip comprising immobilized eosinophil cationic protein (ECP) or other appropriate protein bound to labeled anti-ECP antibody or other specific antibody.

25 In this event, the strip is contacted with mucus from a patient and all of the other parameters, (pH, protein, nitrite, esterase) are read from the strip. The strip is then incubated for a sufficient amount of time so that ECP present in the mucus, due to eosinophil infiltration, competes the labeled antibody from the strip-bound ECP. As a result, upon visualization of the label, the greater the amount of ECP present in the mucus, the lower the amount of label visualized. The antibody could be enzymatically labeled, or labeled with biotin or avidin, which could then be visualized by methods well known in the art. Analogously, ECP or other eosinophil antigen specific antibody could be immobilized on the strip which, upon exposure to nasal secretions, binds any ECP or other eosinophil specific protein or

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enzyme present in the secretion. Excess sample could then be washed from the strip and a second, labeled ECP or other eosinophil antigen specific antibody or chromogenic eosinophil specific enzyme substrate could be contacted with the strip. In this case, the greater the signal upon development, the greater the amount of eosinophilia in the patient.

5 Preferably, an eosinophil specific enzyme is detected by providing a chromogenic reagent on the strip which changes color to an extent proportional to the amount of eosinophils present in the sample. In this context, the teachings of U.S. Patent Nos. 5,369,007 and 5,200,321, herein incorporated by reference, while directed to a very different art (detection of illicit drugs) are instructive here. According to those patents, a microassay on
10 a card could be adapted to the instant utility in either a displacement or competition mode, as described herein above, using eosinophil specific antigens and antibodies. Likewise, the concepts and methods disclosed therein could be utilized to prepare the novel reagent test strip of the instant invention. In yet another embodiment of the novel test strip of this invention, a portion of the strip contacted with the nasal secretion is transparent. This
15 portion is then simply stained and quantitated microscopically for eosinophils as described hereinabove, after reading all of the other reagent parameters.

In any event, even with the use of a standard, commercially available reagent test strip, all that is required for the method of this invention is that, in addition to contacting the nasal
20 secretion with an appropriate reagent test strip and quantitation of the pH, nitrite, esterase and protein, is that the mucus be evaluated for eosinophils. This is quickly and easily achieved by making a smear of the nasal secretion, staining the smear with eosine or other appropriate dye, and quantitating the number of eosinophils present per field. Alternatively, the hemocytometric method of Okuda *et al* (*Int Arch Allergy Immunol.*
25 [1994] 104:6) or of Settipane (*Allergy Proc.* [1985] 6:303-366), herein incorporated by reference, could be used for this purpose.

According to methods known in the art, nasal ECP has been found to range between about 30 ng/ml in a "normal" individual, up to about 200 ng/ml in individuals experiencing acute
30 allergic reactions (see Clin. Exp. Allergy, 1997, 27:270-276; see also JACI 1996, 97:104-112; see also Clin. Exp. Allergy 1994, 24:1151-1156). Commercially available antibodies for carrying out the method and for making the device according to the present invention may be obtained, for example, from Pharmingen. Thus, Pharmingen catalog number 15371A is a mouse IgG1 monoclonal antibody (clone AHE-1) which recognizes human

eosinophil peroxidase, an 81 kD granule protein specific to eosinophils. Pharmingen catalog number 15381A is a mouse IgG1 monoclonal antibody (clone AHE-2) which recognizes human eosinophil Major Basic Protein. a 14 kD granule protein specific to eosinophils. Accordingly, a wide variety of eosinophil specific markers may be employed according to the method and for the device of this invention. Thus, a protein selected from the group, but not limited to the group, eosinophil Major Basic Protein (MBP), present at about 9 µg/million eosinophils, eosinophil cationic protein (ECP), present at about 5 µg/million eosinophils, eosinophil derived neurotoxin (EDN), present at about 3 µg/million eosinophils, or eosinophil peroxidase (EPO), present at about 12 µg/million eosinophils, (see Textbook of Allergy, Principles and Practice, for concentrations of these markers), may be employed in the device or method according to this invention, using commercially available antibody, or antibody developed independently. As mentioned above, avidin/biotin conjugates of such eosinophil specific markers may also be employed according to this invention.

Accordingly, this invention provides a method for rapidly, non-invasively and inexpensively differentiating between allergy and viral or bacterial respiratory infection. The method involves measuring the pH, protein content, nitrite content, esterase activity and eosinophil content of a sample of contacted nasal secretion. A combination of a pH between about 7.5 and 9, a moderately strong presence of protein, and at least a trace of nitrite and esterase and the absence of eosinophils indicates the presence of a bacterial infection. The combination of a pH between about 5.0 and 7.0, little or no protein, little or no nitrite, little or no esterase activity and moderate to strong eosinophil content is indicative of an allergic condition. The combination of a pH between about 5.0 and 7.0, little or no protein, and at least a trace of nitrite and esterase and the absence of eosinophil infiltration indicates an upper respiratory tract viral infection. In a preferred embodiment of this method, each value obtained for pH, protein, nitrite, esterase, and eosinophil infiltration in a patient's nasal secretion, is assigned a value such that the sum of assigned values results in an enhancement in the clustering of patients having bacterial infections, a clustering of patients having viral infection, and a clustering of patients having allergies. At the same time the assigned values preferably result in a separation of these clusters from each other. In a preferred embodiment, this method is practiced with a reagent test strip. The test strips which are used in this testing may be those produced by the Ames division of Miles laboratory as disclosed in any of U.S. Patent Nos. 3,438, 737, 4,301,115;

4,637,979; 4,645,842; 4,657,855; 4,704,460; 4,758,508. or by Boehringer Mannheim Corporation, as disclosed in any of U.S. Patent Nos. 3,359,072; 3,418,079; 3,712,853; 3,897,214; 3,802,842; 4,013,416; 4,385,114. A reagent test strip specifically adapted for rapidly, non-invasively and inexpensively differentiating between allergy and respiratory 5 infection is also provided. This novel strip comprises reagents adapted to provide information on the pH, protein content, nitrite content, esterase activity and eosinophil content of the sample contacted.

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What Is Claimed Is:

1 1. A method for differentiating between allergic rhinitis, upper respiratory tract viral
2 infection and bacterial sinusitis, which comprises measuring a sample of a patient's nasal
3 secretion, for pH, protein content, nitrite content, esterase activity and eosinophil counts of
4 the sample, such that a combination of:

5 (a) a pH between about 7.5 and 9, a moderately strong presence of protein, at least a
6 trace of nitrite and esterase, and low or absent eosinophil counts is indicative of bacterial
7 sinusitis;

8 (b) a pH between about 5.0 and 7.0, little or no protein, little or no nitrite, little or no
9 esterase activity and moderate to high eosinophil counts is indicative of allergic rhinitis;
10 and

11 (c) a pH between about 5.0 and 7.0, little or no protein, at least a trace of nitrite and
12 esterase and low or absent eosinophil counts indicates an upper respiratory tract viral
13 infection.

1 2. The method of claim 1 wherein for each measurement obtained for pH, protein,
2 nitrite, esterase, and eosinophil counts in a patient's nasal secretion, an arbitrary clinical
3 score is assigned such that by summing and graphing the clinical score assigned for each
4 measurement to produce a graph of said clinical scores, a first grouping of clinical scores
5 for patients having bacterial sinusitis occurs, a second grouping of clinical scores for
6 patients having upper respiratory tract viral infection occurs, and a third grouping of
7 clinical scores for patients having allergic rhinitis occurs, such that said graph separates
8 said first, said second and said third grouping of clinical scores.

1 3. The method of claim 1 wherein said method is practiced with a reagent test strip.

1 4. A method for differentiating between viral infectious rhinitis, bacterial sinus
2 infection and allergic rhinitis in a patient, which comprises:

3 (a) contacting nasal secretion from said patient with a reagent test strip adapted for
4 providing measurements of eosinophil, protein, pH, nitrite, and esterase
5 concentrations of the secretion, wherein said reagent test strip is adapted to provide
6 said measurements by providing reactants on said reagent test strip which

- specifically indicate eosinophil, protein, pH, nitrite, and esterase concentrations upon contact of said nasal secretion with said reactants;
- (b) measuring the pH, protein content, nitrite content, esterase activity and eosinophil concentration of said nasal secretion such that a combination of:
- (i) a pH between about 7.5 and 9, a moderately strong presence of protein, at least a trace of nitrite and esterase, and low or absent eosinophil counts is indicative of bacterial sinusitis;
 - (ii) a pH between about 5.0 and 7.0, little or no protein, little or no nitrite, little or no esterase activity and moderate to high eosinophil counts is indicative of allergic rhinitis;
 - (iii) a pH between about 5.0 and 7.0, little or no protein, at least a trace of nitrite and esterase and low or absent eosinophil counts indicates an upper respiratory tract viral infection; and
- (c) correlating the results obtained from said measuring with a control.

5. The method according to claim 1 which comprises contacting and measuring nasal secretion from said patient with a urinalysis reagent test strip, which measures protein, pH, nitrite, and esterase concentrations of the secretion, wherein said reagent test strip is adapted to in addition provide a measurement of the eosinophil concentration of said nasal secretion.

6. The method according to claim 1 which comprises:

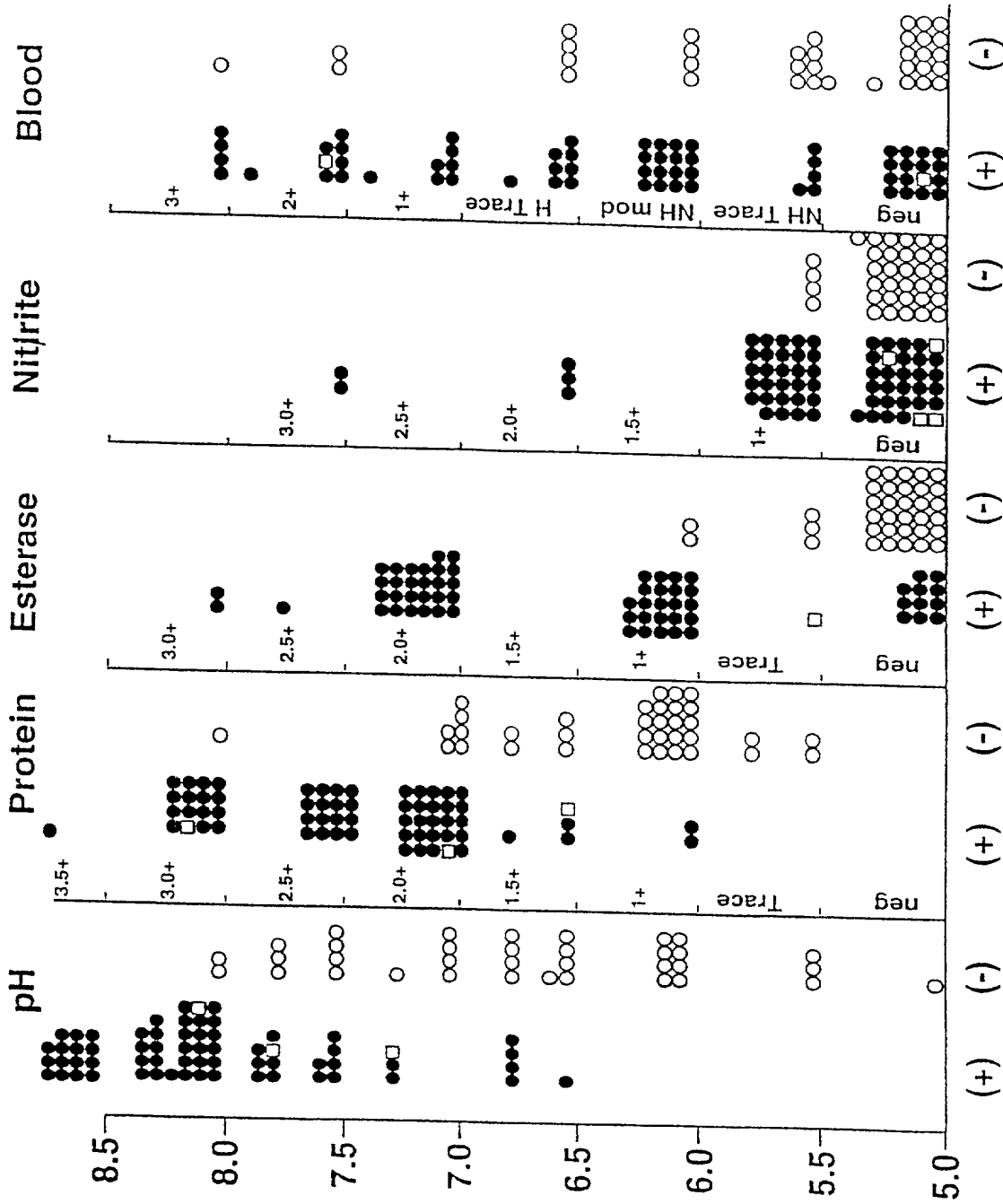
- (a) contacting and measuring nasal secretion from said patient with a urinalysis reagent test strip, which measures protein, pH, nitrite, and esterase concentrations of the secretion; and
- (b) measuring eosinophil concentration of said nasal secretion.

7. A device for differentiating between allergic rhinitis, upper respiratory tract viral infection and bacterial sinusitis, comprising a support upon which is fixed discrete indicators of pH, protein content, nitrite content, esterase activity, and eosinophil content of a sample with which said reagent test strip is contacted.

8. The device according to claim 7 configured as a reagent test strip.

- 1 9. The device according to claim 8 comprising an immobilized eosinophil specific
2 protein.
- 1 10. The device according to claim 9 wherein said protein is selected from the group
2 consisting of eosinophil Major Basic Protein, eosinophil cationic protein, eosinophil
3 derived neurotoxin, eosinophil peroxidase, and mixtures thereof.
- 1 11. The device according to claim 10 wherein said protein is bound to a labeled or unlabeled
2 avidin, biotin, or antibody.
- 1 12. The device according to claim 8 comprising an immobilized antibody specific to an
2 eosinophil specific antigen.
- 1 13. The device according to claim 12 wherein said antibody is specific to a protein selected
2 from the group consisting of eosinophil major basic protein, eosinophil cationic
3 protein, eosinophil derived neurotoxin, eosinophil peroxidase, and mixtures thereof.
- 1 14. The device according to claim 8 comprising an immobilized substrate which upon
2 contact with an eosinophil specific enzyme is converted to a detectable reaction
3 product.
- 1 15. The device according to claim 14 wherein said substrate is a chromogenic substrate.

1/2



Sinusitis
Figure 1

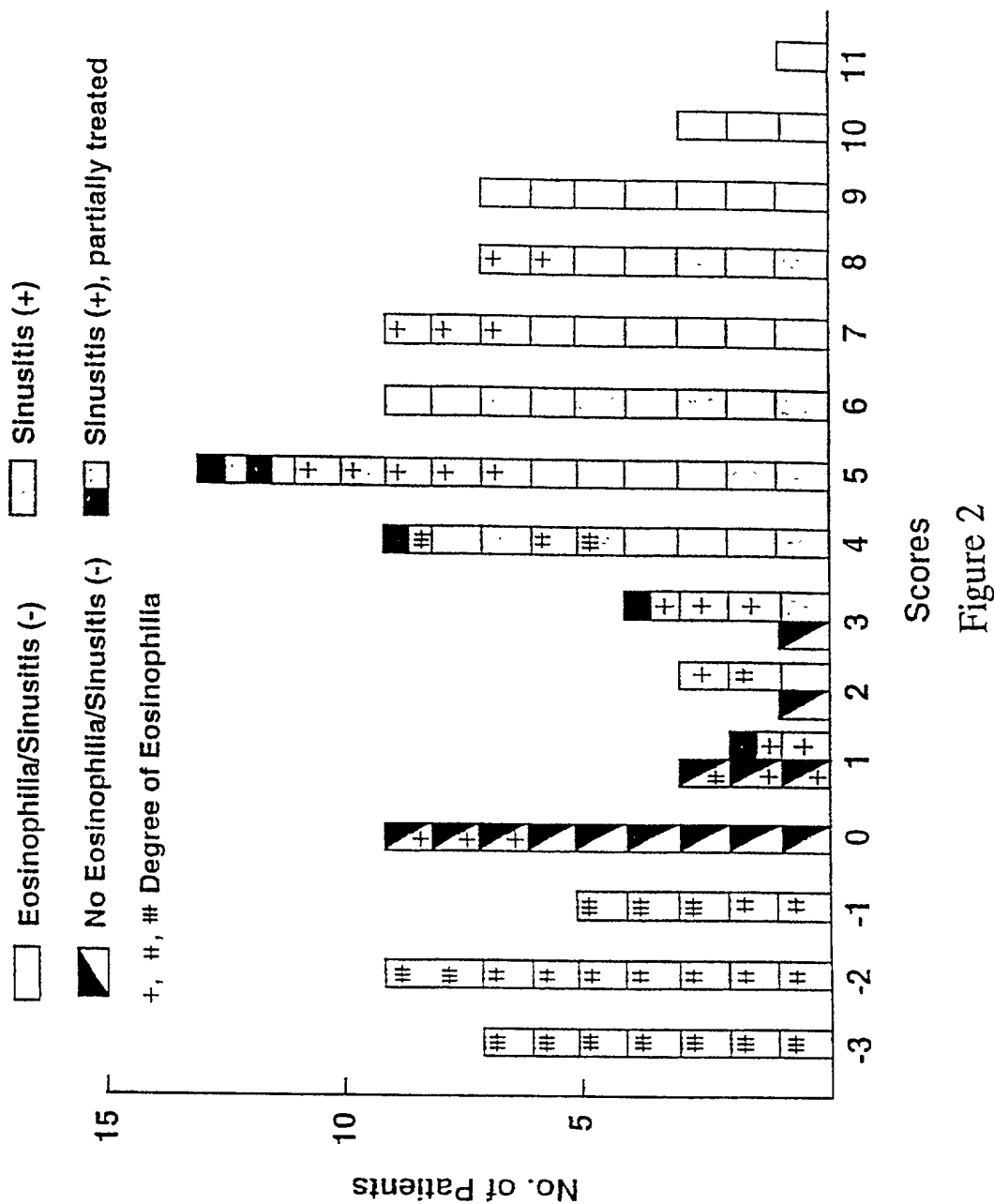


Figure 2

DECLARATION AND POWER OF ATTORNEY
FOR PATENT APPLICATIONATTORNEY DOCKET NO. UF1561A-US

As a below named inventor, I hereby declare that:

My residence/post office address and citizenship are as stated below next to my name;

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled:

Rapid Diagnostic Method for Distinguishing Allergies and Infections

the specification of which is attached hereto unless the following box is checked:

(X) with a filing date of 3/16/1999 as US Application Serial No. 09/936,954 or PCT International Application

Number _____ and was amended on _____ (if applicable).

I hereby state that I have reviewed and understood the contents of the above-identified specification, including the claims, as amended by any amendment(s) referred to above. I acknowledge the duty to disclose all information which is material to patentability as defined in 37 CFR 1.56.

Foreign Application(s) and/or Claim of Foreign Priority

I hereby claim foreign priority benefits under Title 35, United States Code Section 119 of any foreign application(s) for patent or inventor(s) certificate listed below and have also identified below any foreign application for patent or inventor(s) certificate having a filing date before that of the application on which priority is claimed:

COUNTRY	APPLICATION NUMBER	DATE FILED	PRIORITY CLAIMED UNDER 35 U.S.C. 119
PCT	PCT/US99/05751	3/16/1999	YES: ____ NO: ____
			YES: ____ NO: ____

Provisional Application

I hereby claim the benefit under Title 35, United States Code Section 119(e) of any United States provisional application(s) listed below:

APPLICATION SERIAL NUMBER	FILING DATE

U.S. Priority Claim

I hereby claim the benefit under Title 35, United States Code, Section 120 of any United States application(s) listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States application in the manner provided by the first paragraph of Title 35, United States Code Section 112, I acknowledge the duty to disclose material information as defined in Title 37, Code of Federal Regulations, Section 1.56(a) which occurred between the filing date of the prior application and the national or PCT international filing date of this application:

APPLICATION SERIAL NUMBER	FILING DATE	STATUS(patented/pending/abandoned)

POWER OF ATTORNEY:

As a named inventor, I hereby appoint the following attorney(s) and/or agent(s) listed below to prosecute this application and transact all business in the Patent and Trademark Office connected therewith.

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I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

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Inventor's Signature

Date

**DECLARATION AND POWER OF ATTORNEY
FOR PATENT APPLICATION (continued)**

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Inventor's Signature

Date

Full Name of Inventor: _____

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Date